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Twenty-two genetic loci in COPD overlap with population-based lung function and pulmonary fibrosis loci

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Chronic obstructive pulmonary disease (COPD) is a leading cause of mortality worldwide¹. We performed a genetic association in 15,256 cases and 47,936 controls, with replication of select top results ($P < 5 \times 10^{-6}$) in 9,498 cases and 9,748 controls. In the combined meta-analysis, we identified 22 loci at genome-wide significance, of which 15 have been associated with lung function in general population samples, and 4 (*EEFSEC*, *DSP*, *MTCL1*, and *SFTPD*) are novel. We noted 2 loci shared with pulmonary fibrosis (*FAM13A* and *DSP*) but with opposite risk alleles for COPD. None of our loci overlapped with genome-wide associations for asthma; however, one locus has been implicated in the joint susceptibility to asthma and obesity. We also identified genetic correlation between COPD and asthma. Our findings highlight novel loci, demonstrate the importance of specific lung function loci to COPD, and identify potential regions of genetic overlap between COPD and other respiratory diseases.

COPD is characterized by persistent and progressive airflow limitation diagnosed by lung function testing¹. While cigarette smoking is the major risk factor, susceptibility is also influenced by genetics²⁻⁴. We established the International COPD Genetics Consortium (ICGC) to coordinate efforts to find susceptibility loci⁵. We defined cases based on pre-bronchodilator evidence of moderate-to-severe airflow limitation by modified GOLD criteria⁶; controls had normal spirometry, and all analyses were adjusted for age and cigarette smoking (pack-years and smoking status). We performed a two-stage genome-wide association study (Figure 1). In Stage 1, we combined 26 cohorts (Supplemental Table S1) containing 63,192 individuals (15,256 COPD cases and 47,936 controls). We selected 79 loci with $P < 5 \times 10^{-6}$ and in analysis Stage 2, we tested them in the UK BiLEVE dataset (9,498 COPD cases and 9,748 controls) from the UK Biobank and performed an overall meta-analysis (Supplemental Table S3).

We identified 13 genome-wide significant ($P < 5 \times 10^{-8}$) associations in Stage 1. Following the Stage 2 analysis, an additional 9 loci achieved genome-wide significance in the overall meta-

analysis (Table 1 and Figure 2). Of the 22 genome-wide significant loci described in our study, 9 have been previously described as genome- (or exome-) wide significant in studies of COPD^{4,7-10}: *HHIP*, *CHRNA5/15q25*, *HTR4*, *FAM13A*, *RIN3*, *TGFB2*, *GSTCD/NPNT*, *CYP2A6/19q13*, and *16p11.2/IL27*. An additional 8 loci: *ADGRG6/GPR126*, *THSD4*, *ADAM19*, *TET2*, *CFDP1*, *AGER*, *ARMC2*, and *RARB* have been previously described and replicated (Supplemental Table S6) in general population GWASs of two measures of lung function (FEV₁ and FEV₁/FVC) that are used in conjunction to diagnose COPD¹¹⁻¹⁷. One locus near *PID1* was previously associated with FEV₁/FVC, but had not replicated in those studies^{13,17}. Four loci are newly being described as genome-wide significant in association with either COPD or lung function: *EEFSEC*, *DSP*, *MTCL1*, and *SFTPD* (Figure 3).

To explore the potential function and causal genes for our novel loci, in addition to using publicly available datasets and prioritization tools (Supplemental Table S7), we also examined a larger set of lung expression quantitative trait loci (eQTL) in 1038 subjects, including subjects with COPD¹⁸ (Supplemental Table S8). As eQTL are pervasive, we also attempted to determine whether our association signal co-localized¹⁹ with an eQTL signal in lung tissue (Supplemental Table S9). We found strong evidence of co-localization (posterior probability > 0.8) for *DSP*, a major protein of desmosomes required for epidermal integrity²⁰, and *MTCL1*, important in epithelial-cell-specific microtubule stabilization^{21,22}, and expressed in respiratory epithelial cells²³. Variants in strong LD with our top *MTCL1* variant rs647097 appear to have enhancer histone marks in fetal lung fibroblasts^{24,25}. In contrast, we found no evidence of a strong eQTL signal or co-localization at our other two novel loci. At 3q21, *EEFSEC* is a potential candidate, as it is a paralog of *TUFM*, a top blood and lung eQTL gene for the 16p11.2/*IL27* COPD susceptibility locus¹⁰, recently part of a novel COPD-related pathway involving *NLRX1*²⁶⁻²⁸. At 10q22, pulmonary surfactant-associated protein D (*SFTPD*) is the most likely candidate, as it is highly expressed in pneumocytes²³, and *sftpd* (-/-) mice develop pulmonary emphysema²⁹. *SFTPD* has been explored as a COPD biomarker³⁰, and while

rs721917 is not an eQTL, polymorphisms in *SFTPD*, including rs721917, may lead to decreased surfactant protein D levels³¹; though the association of SFTPD polymorphisms with COPD susceptibility have been inconsistent. Our analysis also led to some additional insights into other previously described loci. We found evidence of COPD association and eQTL statistical co-localization in lung tissue (posterior probability > 0.8) for *THSD4*, *HHIP*, *AGER*, *CHRNA3*, and *RARB* (Supplemental Table S9). Additional data on eQTLs (Supplemental Table S8), cohort-specific associations at each locus (Supplemental Figures S1a-v), fine mapping (Supplemental Results and Table S20), and other supportive analysis for previously described and novel loci can be found in the Supplemental Materials.

We noted that our top variant at *DSP* (rs2076295) is also associated ($P = 1.1 \times 10^{-19}$) with pulmonary fibrosis³². Recently, a re-sequencing study³³ at this locus identified a second fibrosis-associated variant (rs2744371). We performed additional analysis to investigate genetic overlap in this region using gwas-pw³⁴ (see Supplement). We found a posterior probability of > 0.99 for overlap at the *FAM13A* locus (top fibrosis SNP, rs2609255; $P_{\text{fibrosis}} = 2.20 \times 10^{-11}$, $P_{\text{COPD}} = 1.9 \times 10^{-7}$) and a posterior probability of 0.84 for overlap near *MAPT/KANSL1* (top $P_{\text{fibrosis}} = 8.87 \times 10^{-14}$, $P_{\text{COPD}} = 4.5 \times 10^{-3}$); while the latter locus did not reach genome-wide significance in our study, we note its independent discovery in a genome-wide association in extremes of lung function¹⁶. Notably, for all four of these variants, the fibrosis risk allele is protective for development of COPD. Emphysema, a key component of COPD, and pulmonary fibrosis are both smoking-related lung diseases that have both shared and distinct pathophysiology³⁵⁻³⁷, though genetic loci with opposing effects have not been previously described. Additional investigation of these loci as well as a more comprehensive assessment of genetic overlap of COPD and pulmonary fibrosis may lead to insight into both disorders.

Because our analysis relied on a spirometric definition of COPD alone, we did not specifically exclude other causes of airway obstruction such as asthma, which can overlap with

COPD in adults³⁸. To define COPD, we used pre-bronchodilator spirometry, which was available across all cohorts, and we included at least moderately affected cases ($FEV_1 < 80\%$ predicted). We examined the top set of genome-wide significant results in a subset of our largest cohorts with both pre- and post-bronchodilator data and densely imputed genotypes; overall, the effect sizes (mean difference = 0.001) and P values (mean \log_{10} P value difference = 0.18) were similar (Supplemental Table S10 and Supplemental Figures S4 and S5). In addition, a recent GWAS of FEV_1 , FVC, and FEV_1/FVC did not find substantial differences including and excluding subjects with asthma¹⁶. In the 79 variants tested in Stage 2, we found no significant difference in the OR for COPD association when including and excluding individuals with asthma (Supplemental Figure S6). We examined COPD associations of genome-wide significant asthma (and asthma-associated traits) loci from the NHGRI-EBI GWAS Catalog³⁹ (Supplemental Table S11). We also compared our COPD association results to the GABRIEL asthma study⁴⁰ (Supplemental Tables S12). None of the genome-wide significant loci from asthma and COPD overlapped. Further, no asthma or COPD loci showed Bonferroni-adjusted (for number of look-ups) significant association with the other disease, though several loci showed nominal ($P < 0.05$) significance (Supplemental Tables S11 and S12). The 16p11.2 (*CCDC101*) locus has been described in the joint susceptibility to asthma and obesity⁴¹. COPD susceptibility is strongly related to cigarette smoking. Two of our loci (15q25 and 19q13) have been previously associated with smoking behavior^{42,43}, though we found no additional evidence of overlap in genome-wide significant variants described in the NHGRI-EBI GWAS Catalog³⁹ and Tobacco and Genetics Consortium GWAS⁴³ (Supplemental Tables S13-S15). In contrast to minimal overlap in genome-wide significant results with asthma and smoking, we discovered a significant overall genetic correlation of COPD with asthma ($r_{\text{genetic}} = 0.38$, $P = 6.2 \times 10^{-5}$) using LD score regression in our white subjects^{44,45}. We also assessed genetic correlation with population-based lung function, pulmonary fibrosis, smoking behavior, and two common COPD comorbidities, coronary artery disease and osteoporosis. We identified significant correlation of COPD with lung

function and two aspects of smoking behavior, but not with common comorbidities or with pulmonary fibrosis (Figure 4). The lack of significant correlation of COPD with pulmonary fibrosis may indicate our overlapping loci for COPD and pulmonary fibrosis are not representative of a broader disease correlation; alternatively, it could reflect limited sample size or a mix of positive and negative genetic correlations across the genome for the diseases. In potential support of this latter hypothesis, and in contrast to the loci we describe in this study, are recent descriptions of rare variants in telomere genes predisposing to both emphysema, a key feature of COPD, and pulmonary fibrosis^{37,46}. Our analysis of partitioned heritability identified COPD genetic association enrichment in fetal lung tissue (coefficient $P = 3.5 \times 10^{-7}$); other analyses also support lung tissue or lung cell types (Supplemental Materials).

Our study is, to our knowledge, the largest genome-wide association study of COPD cases to date and includes over 60,000 subjects (including 15,256 COPD cases) in our Stage 1 analysis. We chose to combine subjects of different ethnicities, hypothesizing that the benefit of shared risk factors across ethnicities would outweigh power loss due to heterogeneity. While methods have been developed that can more rigorously assess the degree of overlap and provide additional power in this setting⁴⁷, none of our non-white cohorts were sufficiently sized or powered for these analyses. COPD is also a highly heterogeneous disease; whether a more precise phenotypic definition would result in greater power is not clear. We used a staged study design and examined overall meta-analysis P-values to determine genome-wide significance. Thus, 9 loci (*TET2*, *CFDP1*, *TGFB2*, *AGER*, *ARMC2*, *PID1*, *MTCL1*, *SFTPD*, and *CYP2A6*) from our Stage 1 analysis, which only reached genome-wide significance in either the Stage 2 UK BiLEVE analysis or the overall meta-analysis, should be further replicated. However, six of these 9 association signals are significant if we consider a Bonferroni correction ($P < 6.3 \times 10^{-4}$) for the 79 variants tested in Stage 2. Further, 8 of these 9 variants become more significant from the Stage 1 analysis to the overall meta-analysis,

with the exception of *RARB*, which has a previously reported association with both lung function¹³ and airflow obstruction¹⁵ (Table 1).

The majority of our significant loci overlap with lung function loci, strengthening the foundation for investigating the relationship of lung function variability in the general population to risk of developing COPD. These loci are unlikely to reflect susceptibility for asthma or for cigarette smoking; however, our association as a whole shows evidence of shared heritability with asthma (supporting investigation into shared genetic etiologies for these diseases) and cigarette smoking behavior (despite adjustment for smoking in our statistical model). We identified enrichment for fetal lung cells, supporting a role for early life events contributing to future risk of COPD. Finally, we identify loci that overlap with pulmonary fibrosis, but with opposite risk alleles. Our study highlights the important contribution of genetic association studies to understanding COPD, not only by identifying novel loci, but also illustrating relationships with other pulmonary traits and diseases.

Online Methods

Study Cohorts

We invited investigators from 22 studies with genome-wide association data and COPD case-control or general population samples with spirometry to participate in a genome-wide association meta-analysis. Additionally, we included four cohorts with Illumina HumanExome v1.2 and custom genotyping based primarily on prior top results from a previously published COPD GWAS⁴, using results with $P < 1 \times 10^{-4}$ using plink '--clump' on the COPDGene non-Hispanic whites to perform linkage disequilibrium pruning ($r^2 < 0.8$), preferentially retaining both an imputed and genotyped top SNP at each locus. An additional group of variants was a candidate panel, based on results from a previous candidate gene analysis⁴⁸, as well as variants identified in association with lung function (supplementing the existing content on the array, which included variants from previous genome-wide association studies), including the lead SNP and a 200kb region around that SNP pruned for variants with $P < 0.01$ and $r^2 < 0.8$, and additional top-ranked SNPs for COPDGene-specific analyses for lung function, bronchodilator responsiveness, exacerbations, and SNPs from candidate genes.

The baseline characteristics of these 26 cohorts can be seen in Supplemental Table S1. Each cohort obtained approval from appropriate ethical/regulatory bodies; informed consent was obtained for all individuals. (Further cohort-specific methods can be found in the Data Supplement.) As most of these cohorts did not have post-bronchodilator spirometry, we used a modified definition of GOLD criteria based on pre-bronchodilator spirometry: forced expiratory volume in 1 second (FEV_1) $< 80\%$ and FEV_1 to forced vital capacity (FVC) ratio of < 0.7 for cases, and $FEV_1 > 80\%$ and $FEV_1/FVC > 0.7$ for controls. Logistic regression was performed in each cohort, adjusting for age, sex, pack-years of smoking, ever-smoking status, current-smoking status, and ancestry-based principal components, as appropriate for each study. Summary statistics were

assessed using EasyQC⁴⁹ version 10.1. More detailed cohort information, including cohort-specific methods, can be found in the Data Supplement.

Genome-wide association quality control

Summary statistics, including effect allele and other allele oriented to the + strand, effect allele frequency, chromosome and position (hg19), and imputation quality were uploaded to a secure site at the Brigham and Women's Hospital / Channing Division of Network Medicine. Quality control assessments included assessing allele frequencies versus 1000 Genomes reference, standard error versus sample size, and quantile-quantile plots. Variants with an imputation quality metric of < 0.3 (provided a higher threshold for imputation quality was not already implemented), a minor allele count (MAC) of < 20 using the effective sample size or the number of cases and adjusted for imputation quality where applicable, were set to missing. Variants were included for meta-analysis if they were present in at least 13 studies (those with European ancestry and at least 7 million markers passing all quality control filters).

Staged GWAS meta-analysis

In Stage 1 of the analysis, we used Metal^{50,51} version 2011-03-25 to perform a fixed-effects meta-analysis of genome-wide data from 22 studies and four additional COPD cohorts genotyped on an Illumina HumanExome v1.2 platform with custom content; this content included a set of COPD candidate genes and regions identified from prior COPD GWAS efforts⁴. We adjusted for inflation using genomic control correction in each study. We included study populations with subjects of non-European ancestry in the overall analysis, and additionally examined results limited to study populations of European ancestry. To identify variants to test for association in Stage 2 in the UK BiLEVE study, we selected top results ($P < 5 \times 10^{-6}$) from the Stage 1 meta-analysis. We selected one lead variant from the chromosome 15q25, *FAM13*, and *HHIP* regions, as all of these have been described in multiple COPD GWASs^{4,7,8,15}. For the remainder of the regions, we performed linkage disequilibrium pruning using the plink2 --clump procedure with an r^2 of 0.5, additionally

examining these SNPs for the number of cohorts with passing quality control at each variant and including SNPs in strong LD (i.e., part of the same clump) with a lower degree of missingness. To identify independent results, we used GCTA-COJO^{52,53} on the Stage 1 meta-analysis for variants with $P < 5 \times 10^{-6}$ using the default distance of 10Mb. We used the COPDGene non-Hispanic whites (as the largest representative population) as the reference population for these analyses. An overall meta-analysis across the Stage 1 and Stage 2 (UK BiLEVE) cohorts was performed and variants with $P < 5 \times 10^{-8}$ were considered genome-wide significant (Figure 1).

Lung eQTL analysis

Lung expression quantitative trait loci (eQTL) were calculated from 1,111 human subjects who underwent lung surgery at three academic sites, Laval University, University of British Columbia (UBC), and University of Groningen, henceforth referred to as Laval, UBC, and Groningen, respectively. This lung eQTL dataset has been described previously^{18,54}. Briefly, 66.7% to 91.2% of the individuals in this study were current or former smokers and 24.2% to 35.3% had moderate to severe COPD (GOLD spirometry grade 2 to 4). Whole-genome gene expression profiling in the lung was performed on a custom Affymetrix array (GPL10379). Microarray pre-processing and quality controls were described previously^{18,55,56}. Probe sequences were mapped to the human genome (hg19) using Bowtie⁵⁷ and probes not mapping to a coding region or having a common SNP (MAF \geq 5%) in their sequence were removed. Expression data were adjusted for age, sex, and smoking status using residuals obtained with the robust fitting of linear models function (rlm) in the R statistical package MASS. Residual values deviating from the median by more than three standard deviations were filtered as outliers. Genotyping was carried on the Illumina Human 1M-Duo BeadChip array.

Twenty-one out of the 22 SNPs (in main manuscript Table 1) were genotyped or imputed in the three cohorts, i.e. Laval, UBC, and Groningen. One of the SNPs, rs7186831, was not well-imputed; a proxy, rs11865296 in modest linkage disequilibrium ($r^2 = 0.54$, 1000 genomes phase 3,

EUR) was used instead. These variants were tested for association with adjusted expression traits (43,465 probe sets) in the lung. SNPs within 1 Mb up and downstream of the transcription probe set were considered as local-eQTL. Distant-acting eQTLs were further than 1 Mb away or on a different chromosome. Association tests were carried with PLINK1.9^{58,59} in each cohort and then meta-analyzed using Fisher's method. All local eQTL with nominal P value < 0.05 in the meta-analysis were considered. To provide an additional overall estimate of eQTL significance, we considered a Bonferroni correction threshold $[(0.05)/(22 \text{ SNPs} \times 43,465 \text{ probe sets}) = P \text{ value} < 5.2 \times 10^{-8}]$. Statistical analyses were performed in R3.2.3⁶⁰.

Co-localization Analysis

Co-localization of statistical signals between COPD genetic association and eQTL were examined using the coloc R package¹⁹. We used phenotypic summary statistics from whites with genome-wide association data and all eQTL results and examined 500kb flanks around the top 22 genome-wide significant associations found in the overall meta-analysis (Table 1).

Sensitivity Analysis

To estimate the effect of using pre- instead of post-bronchodilator lung function on our results, we examined the top set of genome-wide significant results in our largest cohorts with both pre- and post-bronchodilator data and densely imputed genotypes (COPDGene NHW and AA, ECLIPSE, NETT-NAS, and Norway / GenKOLS). Since subjects from these cohorts (except for COPDGene) were included based on post-bronchodilator values, including all subjects with COPD based on post-bronchodilator spirometry would lead to larger sample sizes and make comparison of P-values more difficult. Thus, we chose a random sample of post-bronchodilator cases and controls that matched the number of pre-bronchodilator cases and controls. We performed logistic regression using these equal sized set of pre- and post-bronchodilator cohorts, and meta-analyzed the results.

Asthma overlap analysis

We assessed the overlap between our results and known asthma susceptibility loci. We downloaded information on genome-wide significant ($P < 5 \times 10^{-8}$) associations with asthma and asthma-related traits including asthma and hay fever, asthma (childhood onset), asthma (corticosteroid response), bronchodilator response in asthma, pulmonary function decline, and severe asthma in the NHGRI-EBI GWAS Catalog³⁹. Additionally, we examined top associated variants (which were not genome-wide significant) in the susceptibility to the asthma-COPD overlap syndrome⁶¹. In all, we assessed the association statistics of 49 unique asthma-associated trait loci across 26 genomic regions in our Stage 1 meta-analysis results. We also examined the asthma association statistics of our top COPD loci from overall meta-analysis using publically available asthma GWAS data from the GABRIEL Consortium⁴⁰. For COPD loci not present in the GABRIEL Consortium asthma GWAS data, we attempted to examine proxy SNPs in LD ($r^2 > 0.5$, 1000 genomes phase 1 CEU) with our top COPD loci.

To examine the genetic correlation⁴⁵ of COPD and asthma over the entire genome, we performed LD score regression⁴⁴ using summary statistics from publically available asthma GWAS data from the GABRIEL Consortium⁴⁰. For all comparisons using LD score regression, we filtered to HapMap3 variants, limited to white subjects with genome-wide data, and filtered on missingness using default parameters in `munge_sumstats.py`. For the GABRIEL data, we required a variant to be present in at least 35 of the studies.

Smoking behavior overlap analysis

We downloaded information on genome-wide significant ($P < 5 \times 10^{-8}$) associations with the traits “nicotine dependence” and “smoking behaviour” in the NHGRI-EBI GWAS Catalog³⁹. We assessed these top smoking-associated SNPs in our Stage 1 meta-analysis results. We also assessed overlap of smoking and COPD in the publically available summary statistics from the 2010 Tobacco and Genetics Consortium GWAS⁴³. We evaluated our top COPD loci associations from overall meta-analysis with both cigarettes per day and ever-smoking traits. For COPD risk SNPs not directly

analyzed in the Tobacco and Genetics Consortium GWAS, we attempted to examine proxy SNPs in LD ($r^2 > 0.5$, 1000 genomes phase1 CEU) with our top COPD loci.

To examine the genetic correlation⁴⁵ of COPD and smoking behaviours (cigarettes per day and ever-smoking status) over the entire genome, we performed LD score regression⁴⁴ using summary statistics from our current COPD study as noted above and publically available summary statistics from the 2010 Tobacco and Genetics Consortium GWAS⁴³.

Table 1. Overall study results showing 22 loci with genome-wide significant P values in overall meta-analysis following UK BiLEVE Stage 2 analysis.

rsID	Closest Gene	Locus	Risk Allele	Alt Allele	Risk Allele Mean Freq (Range)	Stage 1 OR (95% CI)	Stage 1 P Value	UK BiLEVE (Stage 2) OR (95% CI)	UK BiLEVE (Stage 2) P Value	Overall Meta-Analysis P Value
rs13141641	<i>HHIP</i>	4q31.21	T	C	0.594 (0.524-0.886)	1.231 (1.183-1.281)	1.16E-24	1.213 (1.16-1.267)	8.15E-18	9.10E-41
rs17486278	<i>CHRNA5</i>	15q25.1	C	A	0.351 (0.244-0.442)	1.224 (1.177-1.273)	2.61E-24	1.126 (1.077-1.178)	2.35E-07	1.77E-28
rs7733088	<i>HTR4</i>	5q32	G	A	0.602 (0.47-0.685)	1.178 (1.129-1.229)	4.40E-14	1.177 (1.127-1.229)	1.78E-13	5.33E-26
rs9399401	<i>ADGRG6</i>	6q24.1	T	C	0.724 (0.615-0.748)	1.141 (1.095-1.189)	3.59E-10	1.174 (1.119-1.232)	6.18E-11	1.81E-19
rs1441358	<i>THSD4</i>	15q23	G	T	0.332 (0.194-0.546)	1.132 (1.09-1.177)	2.06E-10	1.121 (1.071-1.172)	6.87E-07	8.22E-16
rs6837671	<i>FAM13A</i>	4q22.1	G	A	0.405 (0.364-0.582)	1.157 (1.115-1.2)	1.02E-14	1.066 (1.021-1.114)	3.75E-03	7.48E-15
rs11727735	<i>GSTCD</i>	4q24	A	G	0.936 (0.926-0.988)	1.266 (1.167-1.374)	1.55E-08	1.246 (1.144-1.357)	4.93E-07	3.84E-14
rs754388	<i>RIN3</i>	14q32.12	C	G	0.821 (0.804-0.865)	1.196 (1.136-1.258)	7.07E-12	1.108 (1.05-1.169)	1.85E-04	4.96E-14
rs113897301	<i>ADAM19</i>	5q33.3	AT	A	0.175 (0.052-0.187)	1.195 (1.13-1.263)	4.52E-10	1.127 (1.066-1.192)	2.79E-05	1.58E-13
rs2047409	<i>TET2</i>	4q24	A	G	0.618 (0.222-0.649)	1.101 (1.059-1.146)	1.58E-06	1.136 (1.087-1.188)	1.95E-08	2.46E-13*
rs2955083	<i>EEFSEC</i>	3q21.3	A	T	0.881 (0.854-0.893)	1.195 (1.123-1.272)	2.00E-08	1.167 (1.093-1.246)	4.01E-06	4.16E-13
rs7186831	<i>CFDP1</i>	16q23.1	A	G	0.435 (0.226-0.472)	1.125 (1.07-1.183)	3.54E-06	1.116 (1.069-1.165)	6.63E-07	1.12E-11*
rs10429950	<i>TGFB2</i>	1q41	T	C	0.731 (0.216-0.773)	1.117 (1.071-1.164)	1.83E-07	1.096 (1.044-1.15)	1.94E-04	1.66E-10*
rs2070600	<i>AGER</i>	6p21.32	C	T	0.955 (0.85-0.987)	1.276 (1.151-1.414)	3.54E-06	1.208 (1.105-1.319)	2.96E-05	5.94E-10*
rs17707300	<i>CCDC101</i>	16p11.2	C	T	0.373 (0.109-0.433)	1.122 (1.079-1.167)	6.24E-09	1.063 (1.018-1.111)	6.10E-03	6.75E-10
rs2806356	<i>ARMC2</i>	6q21	C	T	0.184 (0.052-0.242)	1.125 (1.071-1.181)	2.84E-06	1.116 (1.057-1.178)	6.88E-05	8.34E-10*
rs16825267	<i>PID1</i>	2q36.3	C	G	0.929 (0.874-0.942)	1.242 (1.149-1.342)	5.22E-08	1.131 (1.045-1.224)	2.27E-03	1.68E-09*
rs2076295	<i>DSP</i>	6p24.3	T	G	0.554 (0.442-0.581)	1.107 (1.067-1.149)	4.95E-08	1.061 (1.016-1.107)	7.45E-03	3.97E-09
rs647097	<i>MTCL1</i>	18p11.22	C	T	0.269 (0.259-0.399)	1.106 (1.06-1.154)	3.03E-06	1.089 (1.038-1.142)	4.66E-04	6.14E-09*
rs1529672	<i>RARB</i>	3p24.2	C	A	0.831 (0.675-0.862)	1.162 (1.106-1.221)	2.37E-09	1.048 (0.991-1.109)	9.95E-02	2.47E-08
rs721917	<i>SFTPD</i>	10q22.3	G	A	0.422 (0.392-0.631)	1.092 (1.053-1.132)	2.11E-06	1.068 (1.023-1.115)	2.60E-03	2.49E-08*
rs12459249	<i>CYP2A6</i>	19q13.2	C	T	0.662 (0.617-0.702)	1.126 (1.071-1.183)	2.89E-06	1.077 (1.029-1.127)	1.35E-03	3.42E-08*

OR = odds ratio, CI = confidence interval. *Genome-wide significant in overall meta-analysis only

Figure 1. Study design showing cohorts used in each stage of the analysis.

Stage 1:

26 studies represented. European ancestry studies are shaded.

Study	COPD Cases	Controls
ARIC	1060	6164
B58	205	3665
CHS EA	736	1586
COPACETIC	397	1906
COPDGene NHW	3068	2110
ECLIPSE	1741	149
EOCOPD*	394	495
ICGN*	1852	557
EQTL	252	224
FHS	701	5110
LifeLines	466	9863
Lovelace	259	641
MESA Caucasian	167	754
NETT-NAS	376	435
Norway/GenKOLS	846	695
RS1	112	815
RS2	94	811
RS3	106	1596
SPIROMICS	571	175
TCGS-Poland *	307	311
CHS AA	138	292
COPDGene AA	910	1556
KARE	199	6741
MESA AA	94	532
MESA Hispanic	52	548
TCGS-Korea *	153	205
TOTAL	15256	47936

Stage 2:

Top results from Stage 1 analysis tested in UK BiLEVE study.

Study	COPD Cases	Controls
UK BiLEVE Never Smokers	3737	4871
UK BiLEVE Heavy Smokers	5761	4877
TOTAL	9498	9748

ARIC = Atherosclerosis Risk in Communities, B58 = British 1958 Birth Cohort, CHS = Cardiovascular Health Study, COPACETIC = COPD Pathology: Addressing Critical gaps, Early Treatment & Diagnosis and Innovative Concepts, ECLIPSE = Evaluation of COPD Longitudinally to Identify Predictive Surrogate End-points, eQTL = Lung Expression Quantitative Trait Loci Study, FHS = Framingham Heart Study, KARE = Korean Association Resource project, MESA = Multi-Ethnic Study of Atherosclerosis, NETT-NAS = National Emphysema Treatment Trial / Normative Aging Study, RS = Rotterdam Study, SPIROMICS = Subpopulations and intermediate outcome measures in COPD study, EOCOPD = Boston Early-Onset COPD Study, ICGN = International COPD Genetics Network, TCGS = Transcontinental COPD Genetics Study, UK BiLEVE = UK Biobank Lung Exome Variant Evaluation; NHW = Non-Hispanic white, AA = African American, EA = European American. * Studies without genome-wide array genotyping (custom genotyping)

Figure 2. P values for Stage 1 analysis (small open diamonds) with overlay of overall meta-analysis P values for SNPs analyzed in UK BiLEVE Stage 2 analysis (filled circles). Gene names in gray are previously described COPD or lung function (FEV₁ or FEV₁/FVC) loci; black are novel loci discovered in this study. The Stage 1 cohorts with available genotyping data (Supplemental Figures S1a-v) and the UK BiLEVE cohort determined the sample size for each top variant.

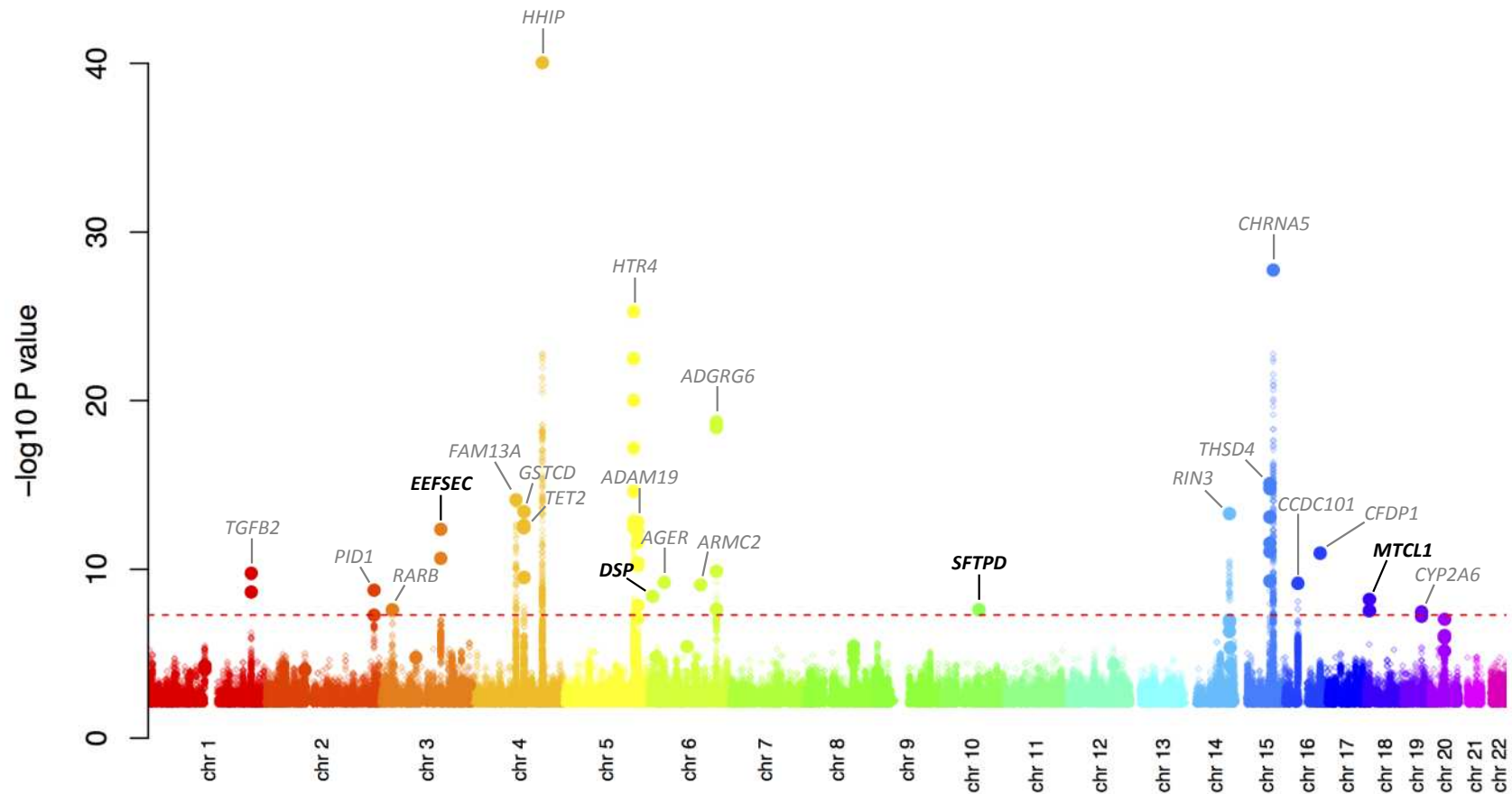


Figure 3a-d. Regional association for novel loci. LocusZoom plots showing regional association of variants at the four novel COPD loci. The point size is proportional to the sample size, where Stage 1 cohorts with available genotyping data (Supplemental Figures S1a-v) and the UK BiLEVE cohort determined the sample size for each top variant.

Fig 3a. rs2955083 (3q21.3, *EEFSEC*)

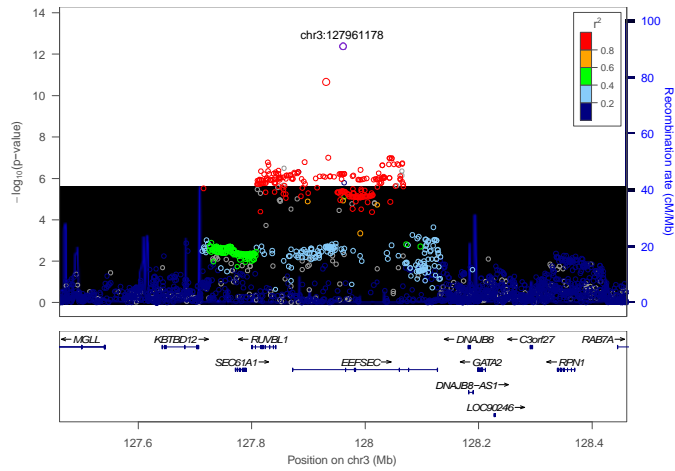


Fig 3b. rs2076295 (6p24.3, *DSP*)

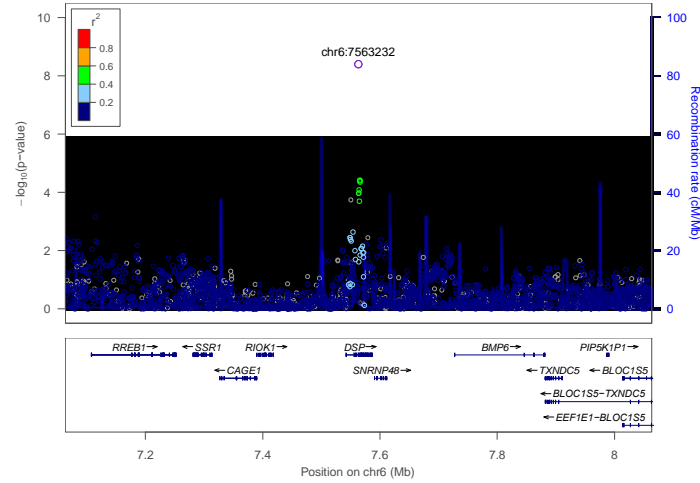


Fig 3c. rs647097 (18p11.22, *MTCL1*)

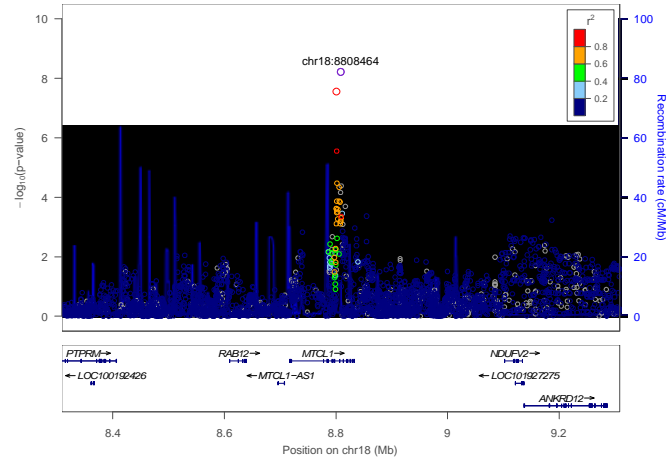


Fig 3d. rs721917 (10q22.3, *SFTPD*)

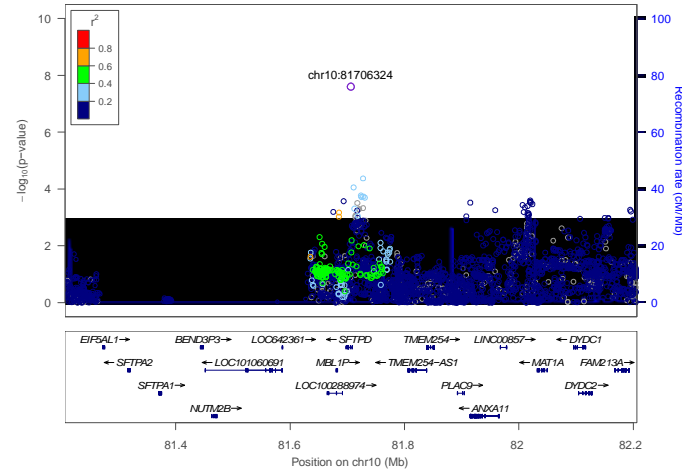
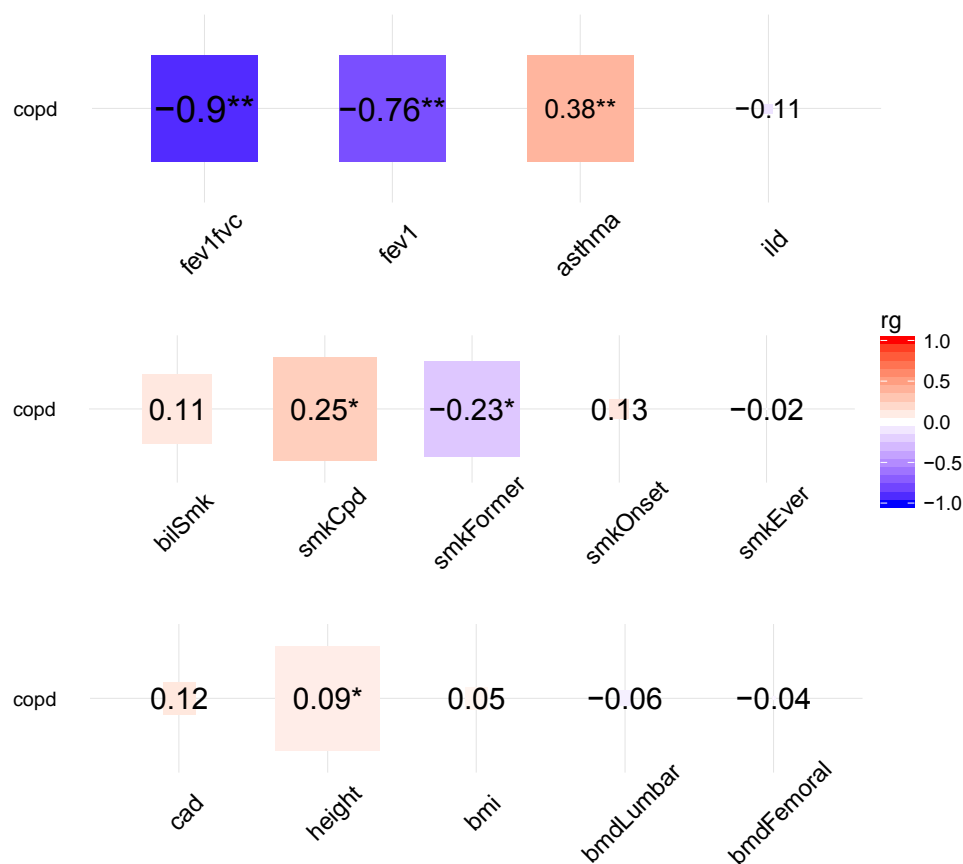


Figure 4 Genetic correlation (using LD score regression) between COPD and other traits.

Shading and numbers represents strength of correlation. * indicates nominal ($P < 0.05$) significance, ** indicates significant after Bonferroni correction for number of pairwise comparisons. fev1fvc and fev1 = lung function (FEV_1/FVC ratio and FEV_1 from CHARGE/SpiroMeta¹³, asthma taken from the asthma GWAS by the GABRIEL Consortium⁴⁰, ild = pulmonary fibrosis from Fingerlin et al.^{32,62}, bilSmk = subset of smokers in the UK BiLEVE study¹⁶, smkCpd = cigarettes per day smoking from the Tobacco and Genetics (TAG) Consortium⁴³, smkFormer = current versus former smokers from TAG, smkOnset = age of smoking initiation from TAG, smkEver = ever versus never smoking from TAG. cad = coronary artery disease from the CARDIoGRAM study⁶³, height⁶⁴ and bmi (body mass index)⁶⁵ from the GIANT consortium, bmdLumbar and bmdFemoral = lumbar and femoral bone mineral density, respectively, from the Genetic Factors for Osteoporosis (GeFOS) Consortium⁶⁶.



Author Contributions:

B.D.H. & M.H.C. contributed to the study concept and design, data analysis, statistical support, and manuscript writing. K.d.J., S.J.L., & D.P.S. contributed to the study concept and design & data analysis. N.S. & M.S.A. contributed to the data analysis and statistical support. T.H.B. & J.E.H. contributed to the study concept and design and statistical support. L.L. contributed to the data collection, data analysis, and statistical support. K.E.N. contributed to data collection and data analysis. J.D.C., B.M.P., R.T.S., G.T.O., Y.T., R.G.B., S.I.R., P.B., A.G., P.G.W., D.A.M., D.A.S., & E.K.S. contributed to the study concept and design and data collection. D.Q., T.A.F., M.L., Y.B., N.S., A.B.W., N.F., P.J.C., R.P.C., T.M.B., S.A.G., J.C.L., J.D., J.B.W., M.K.L., S.L., A.M., X.W., & E.J.A. contributed to the data analysis. L.V.W., I.P.H., P.D.P., D.S.P., W.M., M.D.T., & H.M.B. contributed to the study concept and design. S.R.H., P.A.D., W.J.K., Y.O., S.S.R., D.S., A.L.L., G.G.B., B.H.S., E.R.B., D.A.L., J.J.Y., D.K.K., I.H., P.S. & M.H. contributed to the data collection. All authors, including those whose initials are not listed above, contributed to the critical review and editing of the manuscript and approved the final version of the manuscript.

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N.L. and R.T-S. are shareholders and employees of GSK.

S.I.R. is a current employee and shareholder at AstraZeneca. He has served as a consultant, participated in advisory boards, received honorarium for speaking or grant support from: American Board of Internal Medicine, Advantage Healthcare, Almirall, American Thoracic Society, AstraZeneca, Baxter, Boehringer Ingelheim, Chiesi, ClearView Healthcare, Cleveland Clinic, Complete Medical Group, CSL, Dailchi Sankyo, Decision Resources, Forest, Gerson Lehman, Grifols, GroupH, Guidepoint Global, Haymarket, Huron Consulting, Inthought, Johnson and Johnson, Methodist Health System – Dallas, NCI Consulting, Novartis, Pearl, Penn Technology, Pfizer, PlanningShop, PSL FirstWord, Qwessential, Takeda, Theron and WebMD.

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